	Application No.	Applicant(s)
	00/990 039	DIGHADDOFTAL
Notice of Allowability	09/889,938 Examiner	RICHARDS ET AL. Art Unit
	Ashwin Mehta	1638
The MAILING DATE of this communication appear All claims being allowable, PROSECUTION ON THE MERITS IS herewith (or previously mailed), a Notice of Allowance (PTOL-85) NOTICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RI	(OR REMAINS) CLOSED in or other appropriate commu IGHTS. This application is s	this application. If not included unication will be mailed in due course. THIS
1. This communication is responsive to papers filed 27 Augus	st 2004.	
2. The allowed claim(s) is/are <u>62-65,67,68 and 70-72</u> .		
3. \boxtimes The drawings filed on <u>11 December 2001</u> are accepted by	the Examiner.	
 4. Acknowledgment is made of a claim for foreign priority una) All b) Some* c) None of the: Certified copies of the priority documents have Certified copies of the priority documents have Copies of the certified copies of the priority documents have Copies of the certified copies of the priority documents have linternational Bureau (PCT Rule 17.2(a)). * Certified copies not received: 	been received. been received in Applicatio	n No
Applicant has THREE MONTHS FROM THE "MAILING DATE" noted below. Failure to timely comply will result in ABANDONM THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.		a reply complying with the requirements
5. A SUBSTITUTE OATH OR DECLARATION must be submit INFORMAL PATENT APPLICATION (PTO-152) which give	itted. Note the attached EXA es reason(s) why the oath or	MINER'S AMENDMENT or NOTICE OF declaration is deficient.
6. CORRECTED DRAWINGS (as "replacement sheets") mus	t be submitted.	
(a) ☐ including changes required by the Notice of Draftspers		(PTO-948) attached
1) hereto or 2) to Paper No./Mail Date	<u>-</u>	,
(b) including changes required by the attached Examiner's Paper No./Mail Date	s Amendment / Comment or	in the Office action of
Identifying indicia such as the application number (see 37 CFR 1. each sheet. Replacement sheet(s) should be labeled as such in the	84(c)) should be written on th	e drawings in the front (not the back) of
7. DEPOSIT OF and/or INFORMATION about the deposit attached Examiner's comment regarding REQUIREMENT I	sit of BIOLOGICAL MATE	RIAL must be submitted. Note the
Attachment(s) 1. Notice of References Cited (PTO-892)	5. ☐ Notice of Inf	ormal Patent Application (PTO-152)
2. Notice of Draftperson's Patent Drawing Review (PTO-948)		mmary (PTO-413),
3. Information Disclosure Statements (PTO-1449 or PTO/SB/0	Paper No./i	Mail Date <u>attached</u> . Amendment/Comment
Paper No./Mail Date4. Examiner's Comment Regarding Requirement for Deposit	8 M Evaminada	Statement of Reasons for Allowance
of Biological Material	9. Other	

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Claim Objections and Rejections

- 1. The rejection of claims 62-64, 66, and 69-72 under 35 U.S.C. 112, 2nd paragraph, is withdrawn in light of the claim amendments.
- 2. The rejection of claims 62-72 under 35 U.S.C. 103(a) is withdrawn in light of the claim amendments below and further consideration.

Examiner's Amendment

3. An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Gwen Wood on September 30, 2004.

The application has been amended as follows:

In the claims:

Claim 62. A method for conveying resistance to beet necrotic yellow vein virus (BNYVV) to a sugar beet plant, comprising:

preparing a <u>BNYVV</u> DNA fragment <u>consisting of a nucleotide sequence</u> that corresponds to nucleotides 153 to 3258 of RNA1 of said virus[, wherein said nucleotides 153 to 3258 of RNA1 represent a 3' truncated sequence of BNYVV, and wherein a 5' end primer and a 3' end

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primer are used to obtain the DNA fragment, said 5' end primer consisting of 5'-

CGCGGATCCACCATGGCAGATTCGTTC-3' (containing a BamHI and NcoI restriction site and nucleotides identical to nucleotides 153-168) or 5'-GACGAATTCAAGTCGTCTTTC-3' (containing an EcoRI restriction site and nucleotides complementary to nucleotides 288-301); and said 3' end primer consisting of 5'-GACGAATTCGAAAGATGAGTCTA-3' (containing an EcoRI site and nucleotides identical to nucleotides 2799-2812) or 5'-

CGCAGATCTTTAACTGCTCATCACCAAC-3' (containing a BglII site and nucleotides complementary to nucleotides 3244-3258); and stop codon];

introducing said DNA fragment, operatively linked to a promoter that is active in sugar beet plants, into a sugar beet plant cell to obtain a transformed sugar beet cell; and regenerating a transgenic sugar beet plant from the transformed sugar beet plant cell, said transformed sugar beet plant cell comprising in its genome at least two copies of the DNA fragment, wherein expression of said DNA fragment conveys resistance to BNYVV to said transgenic sugar beet plant.

Claim 64. A transformation vector for conveying resistance to BNYVV to a plant, comprising a BNYVV DNA fragment consisting of a nucleotide sequence [DNA fragment] that corresponds to nucleotides 153 to 3258 of RNA1 of said virus, and transcription and translation regulatory sequences operably linked therewith[, wherein said nucleotides 153 to 3258 of RNA1 represent a 3' truncated sequence of BNYVV, and wherein a 5' end primer and a 3' end primer are used to obtain the DNA fragment, said 5' end primer consisting of 5'-

CGCGGATCCACCATGGCAGATTCGTTC-3' (containing a BamHI and NcoI restriction site

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and nucleotides identical to nucleotides 153-168) or 5'-GACGAATTCAAGTCGTCTTTC-3' (containing an EcoRI restriction site and nucleotides complementary to nucleotides 288-301); and said 3' end primer consisting of 5'-GACGAATTCGAAAGATGAGTCTA-3' (containing an EcoRI site and nucleotides identical to nucleotides 2799-2812) or 5'-CGCAGATCTTAACTGCTCATCACCAAC-3' (containing a BglII site and nucleotides complementary to nucleotides 3244-3258); and stop codon].

Claim 65. A transgenic <u>sugar beet</u> plant cell, exhibiting resistance to BNYVV, comprising in its genome at least two copies of a <u>BNYVV</u> DNA fragment <u>consisting of a nucleotide sequence that corresponds to nucleotides 153 to 3258 of [comprising]</u> RNA1 of said virus[, wherein a 5' end primer and a 3' end primer are used to obtain the DNA fragment, said 5' end primer consisting of 5'-CGCGGATCCACCATGGCAGATTCGTTC-3' (containing a BamHI and NcoI restriction site and nucleotides identical to nucleotides 153-168) or 5'-GACGAATTCAAGTCGTCTTTC-3' (containing an EcoRI restriction site and nucleotides complementary to nucleotides 288-301); and said 3' end primer consisting of 5'-GACGAATTCGAAAGATGAGTCTA-3' (containing an EcoRI site and nucleotides identical to nucleotides 2799-2812) or 5'CGCAGATCTTTAACTGCTCATCACCAAC-3' (containing a BglII site and nucleotides complementary to nucleotides 3244-3258); and stop codon].

Claim 66 was cancelled.

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Claim 68. A transgenic sugar beet plant exhibiting resistance to BNYVV, comprising plant cells having in their genome at least two copies of a BNYVV DNA fragment consisting of a nucleotide sequence that corresponds to nucleotides 153 to 3258 of RNA1 of said virus, and transcription and translation regulatory sequences operably linked therewith [comprising RNA1 of said virus, wherein a 5' end primer and a 3' end primer are used to obtain the DNA fragment, said 5' end primer consisting of 5'-CGCGGATCCACCATGGCAGATTCGTTC-3' (containing a BamHI and NcoI restriction site and nucleotides identical to nucleotides 153-168) or 5'-GACGAATTCAAGTCGTCTTTC-3' (containing an EcoRI restriction site and nucleotides complementary to nucleotides 288-301); and said 3' end primer consisting of 5'-GACGAATTCGAAAGATGAGTCTA-3' (containing an EcoRI site and nucleotides identical to nucleotides 2799-2812) or 5'-CGCAGATCTTTAACTGCTCATCACCAAC-3' (containing a BgIII site and nucleotides complementary to nucleotides 3244-3258); and stop codonl.

Claim 69 was cancelled.

In claim 70, line 4, the recitation, "comprising RNA1 of said virus" was deleted.

In claim 71, line 1, the recitation, "The seeds" was replaced with --Seeds--; and in line 4, the recitation, "comprising RNA1 of said virus" was deleted.

In claim 72, line 1, the recitation, "The vegetatively" was replaced with -- Vegetatively--.

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In the abstract:

The abstract was amended as follows:

The present invention relates to a method for conveying resistance to beet necrotic yellow vein virus (BNYVV) to a sugar beet plant, which method comprises the following steps: (a) preparing [nucleotides in a sequence that is essentially] a DNA fragment consisting of a nucleotide sequence that corresponds to nucleotides 153 to 3258 [of at least 15 nucleotides homologous to the corresponding nucleotide sequence] of the genomic RNA 1 of the beet necrotic yellow vein virus (BNYVV); (b) introducing said DNA fragment, operatively linked to a promoter that is active in sugar beet plants, into a sugar beet plant cell to obtain a transformed sugar beet cell; and (c) regenerating a transgenic sugar beet plant from the transformed sugar beet plant cell.

- 4. Claims 62-65, 67, 68, and 70-72 are allowed.
- 5. The following is an examiner's statement of reasons for allowance: Applicants have developed a method to confer resistance to sugar beet plants against beet necrotic yellow vein virus (BNYVV). The method comprises transforming sugar beet plant cells with a construct comprising a promoter operably linked to a DNA fragment consisting of a nucleotide sequence that corresponds to nucleotides 153-3258 of RNA1 of BNYVV, and regenerating transgenic plants from the transformed cells. The specification teaches that transgenic plants comprising a

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single copy of the introduced DNA remained susceptible to BNYVV, whereas transgenic plants comprising two or more copies showed resistance. Nucleotides 153-3258 are from the portion of BNYVV RNA1 that encodes the viral replicase. Baulcombe (Plant Cell, 1996, Vol. 8, pages 1833-1844) discusses the successful expression of viral replicase genes and portions of viral replicase genes in transgenic plants to confer resistance against the virus from which the gene was derived. In some examples, the resistance was conferred through RNA-mediated homology-dependent resistance, and in other examples the resistance was protein-based. However, while the prior art discusses the expression of sequences from plant viral genes in transgenic plants, to confer resistance against that virus, the prior art does not teach or fairly suggest the expression of specifically nucleotides 153-3258 of RNA1 of BNYVV to confer viral resistance.

Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

Contact Information

Any inquiry concerning this or earlier communications from the Examiner should be directed to Ashwin Mehta, whose telephone number is 571-272-0803. The Examiner can normally be reached from 8:00 A.M to 5:30 P.M. If attempts to reach the Examiner by telephone are unsuccessful, the Examiner's supervisor, Amy Nelson, can be reached at 571-272-0804. The fax phone numbers for the organization where this application or proceeding is assigned are 703-872-9306 for regular communications and 703-872-9307 for After Final communications. Patent applicants with problems or questions regarding electronic images that can be viewed in the Patent Application Information Retrieval system (PAIR) can now contact the USPTO's Patent Electronic Business Center (Patent EBC) for assistance. Representatives are available to answer your questions daily from 6 am to midnight (EST). The toll free number is (866) 217-9197. When calling please have your application serial or patent number, the type of document you are having an image problem with, the number of pages and the specific nature of the problem. The Patent Electronic Business Center will notify applicants of the resolution of the problem within

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October 1, 2004

Ashwin D. Mehta, Ph.D. Primary Examiner

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